Amendment to the Claims:

Please amend the claims as shown in the following listing of claims:

Claims 1-26 (canceled)

- 27. (withdrawn currently amended) A method of preparing a sialyl Lewis x determinant, the method comprising contacting a truneated recombinant murine Fuc-TVII enzyme with a GDP-fucose donor substrate and a sialyl-N-acetyl-lactosamine acceptor substrate in an enzyme bioreactor under conditions that allow the addition of an α1,3 linked fucose to the sialyl-N-acetyl-lactosamine acceptor substrate, wherein the truneated recombinant murine Fuc-TVII enzyme comprises a catalytic domain that is encoded by a nucleic acid sequence segment that is identical to a polynucleotide that is amplified using murine mRNA or cDNA as a template by a 5' primer as shown in SEQ ID NO:3 and a 3' primer as shown in SEQ ID NO:4.
- (withdrawn) The method of claim 27, wherein the Fuc-TVII enzyme is in solution.
- (withdrawn) The method of claim 27, wherein the Fuc-TVII enzyme is immobilized on a solid phase matrix.
- (withdrawn) The method of claim 27, wherein the Fuc-TVII enzyme is a recombinant enzyme.
- (withdrawn) The method of claim 20, wherein the Fuc-TVII enzyme is produced in a mammalian host cell.
- (withdrawn) The method of claim 20, wherein the Fuc-TVII enzyme is produced in a baculovirus host.
- (withdrawn) The method of claim 27, wherein the sialyl-N-acetyl-lactosamine acceptor is on a glycoprotein.
- (withdrawn) The method of claim 27, wherein the sialyl-N-acetyl-lactosamine acceptor is on a glycolipid.
- (withdrawn) The method of claim 27, wherein the sialyl-N-acetyl-lactosamine acceptor is a free oligosaccharide.

- 36. (canceled)
- 37. (currently amended) A truneated recombinant murine Fuc-TVII enzyme comprising a catalytic domain that is encoded by a nucleic acid sequence segment that is identical to a polynucleotide that is amplified using murine mRNA or cDNA as a template by a 5' primer as shown in SEQ ID NO:3 and a 3' primer as shown in SEQ ID NO:4.
- (currently amended) The <u>recombinant</u> murine Fue-TVII enzyme of claim 37, wherein the catalytic domain is encoded by a nucleic acid segment consisting of residue 2194 to residue 3085 of SEO ID NO:1.